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## 2p15p16.1 microdeletion syndrome

**INSERM** 

## Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>2p15p16.1</u> <u>microdeletion syndrome</u>. ORPHA:261349

2p15p16.1 microdeletion syndrome is a recently described syndrome characterized by developmental delay and facial dysmorphism.

Qeios ID: 0MX20V · https://doi.org/10.32388/0MX20V